

AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1-20. (cancelled)

21. (currently amended): A method for preparing monodisperse biodegradable microspheres comprising the steps of:

a) preparing an emulsion comprising at least one polymer organic phase, which comprises an active ingredient and a biodegradable polymer dissolved in an organic solvent, and at least one aqueous phase, the viscosity of the organic phase and the aqueous phase having a ratio of from 0.1 to 10;

b) subjecting the emulsion obtained to controlled laminar shearing;

c) removing the solvent from the polymer organic phase; and

d) isolating the microspheres so obtained.

22. (currently amended): The method of claim 21, wherein the majority of the microspheres are constituted in majority by [[a]] the biodegradable polymer.

23. (previously presented): The method of claim 22, wherein the biodegradable polymer is selected from poly(α -hydroxy) acids, the aliphatic polyesters of poly(α -hydroxy acids), of poly(ϵ -caprolactones)-PCL, of polydioxanones - PDO, polyorthoesters, polyanhydrides, polycyanoacrylates, polyurethanes, polypeptides or poly(amino acids), modified polysaccharides, cellulose, polycarbonates, polydimethylsiloxanes and poly(vinyl acetates) and their derivatives and copolymers.

24. (previously presented): The method of claim 22, wherein the biodegradable polymer is selected from polylactic acids (PLA), and the copolymers of polylactic acid / polyglycolic acid (PLGA).

25. (currently amended): The method of claim 21, wherein the biodegradable polymer has a molecular weight of from 50 to 500 kDaltons.

26. (previously presented): The method of claim 21, wherein the organic solvent of the organic phase of the emulsion is ethyl acetate.

27. (previously presented): The method of claim 21, wherein the active ingredient is lipid-soluble.

28. (previously presented): The method of claim 21, wherein the active ingredient is water-soluble.

29. (previously presented): The method of claim 21, wherein the active ingredient is a peptide or a protein.

30. (previously presented): The method of claim 21, wherein the emulsion prepared in step (a) comprises a hydrophilic active ingredient in combination with a lipophilic active ingredient.

31. (previously presented): The method of claim 21, wherein the organic phase of the emulsion represents from 10 to 60% by weight relative to the total weight of the emulsion.

32. (currently amended): The method of claim 21, wherein the organic phase of the emulsion comprises from 1 to 50%, ~~preferably from 5 to 30% by weight of polymer.~~

33. (currently amended): The method of claim 21, wherein the organic phase of the emulsion comprises from 1 to 50%, ~~preferably from 5 to 30% by weight of active ingredient.~~

34. (previously presented): The method of claim 21, wherein the emulsion is a double emulsion.

35. (previously presented): The method of claim 21, wherein the external and/or internal aqueous phase of the emulsion contains at least one stabilizing agent and/or at least one viscosity agent.

36. (previously presented): The method of claim 21, wherein the external and/or internal aqueous phase of the emulsion contains at least one stabilizing agent and/or at least one osmolarity agent and/or at least one surfactant and/or at least one buffer agent.

37. (previously presented): The method of claim 21, wherein the step of calibration by laminar shearing is carried out in a Couette device.

38. (currently amended): The method of claim 21, wherein the step of removing the solvent from the polymer organic phase is carried out by extraction in water.

39. (currently amended): A method for the administration of active ingredients in the human or animal organism, comprising making use of administering the microspheres ~~that can be obtained~~ comprising an active ingredient and being prepared by the method according to claim 21.

40. (previously presented): The method of claim 39, wherein the active ingredient is selected from antibiotics, hypolipidaemics, antihypertensives, antiviral agents, beta blockers, bronchodilators, cytostatics, psychotropic agents, hormones, vasodilators, anti-allergics, analgesics, antipyretics, antispasmodics, anti-inflammatories, anti-angiogenics, antibacterials, anti-ulcerants, antifungals, anti-parasitics, antidiabetics, anti-epileptics, anti-Parkinsons, antimigraines, anti-Alzheimers, anti-acneics, antiglaucomic agents, anti-asthmatics, neuroleptics, antidepressants, anxiolytics, hypnotics, normothymics, sedatives, psychostimulants, anti-osteoporosis agents, anti-arthritis, anticoagulants, antipsoriasis agents, hyperglycaemics, orexigenics, anorexigenics, anti-asthenics, anticonstipation agents, antidiarrhoeals, anti-trauma agents, diuretics, myorelaxants, enuresis medicaments, erection disorder medicaments, vitamins, peptides, proteins, anticancer agents, nucleic acids, RNA, oligonucleotides, ribozymes and DNA.